existing stocks of products, containing resmethrin until supplies are exhausted, provided that such sale, distribution, or use is consistent with the terms of the previously approved labeling on, or that accompanied, the canceled products.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: June 18, 2013.

Michael Goodis,

Acting Director, Pesticide Re-Evaluation Division, Office of Pesticide Programs.

[FR Doc. 2013–15320 Filed 6–25–13; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2013–0401; FRL–9390–7]

Registration Review; Pesticide Dockets Opened for Review and Comment; Announcement of Registration Review Case Closures

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: With this document, EPA is opening the public comment period for several registration reviews. Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. Registration review dockets contain information that will assist the public in understanding the types of information and issues that the Agency may consider during the course of registration reviews. Through this program, EPA is ensuring that each pesticide's registration is based on current scientific and other knowledge, including its effects on human health and the environment. This document also announces the registration review case closures for the pesticides oxydemeton-methyl (ODM) (case #0258) and resmethrin (case #0421), and the availability of their respective Case Closure Documents. The cancellation of all ODM registrations will become effective on December 31, 2014. The cancellation of all resmethrin registrations will become effective December 31, 2015. These case closures are being announced herein with no comment period.

DATES: Comments must be received on or before August 26, 2013.

ADDRESSES: Submit your comments identified by the docket identification (ID) number for the specific pesticide of interest provided in the table in Unit III.A., by one of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the online instructions for submitting comments. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute.
- *Mail:* OPP Docket, Environmental Protection Agency Docket Center (EPA/DC), (28221T), 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001.
- *Hand Delivery:* To make special arrangements for hand delivery or delivery of boxed information, please follow the instructions at *http://www.epa.gov/dockets/contacts.htm.*

Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is available at *http://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: *For pesticide specific information contact:* The Chemical Review Manager identified in the table in Unit III.A. for the pesticide of interest.

For general information contact: Kevin Costello, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 305–5026; fax number: (703) 308–8090; email address: *costello.kevin@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

This action is directed to the public in general, and may be of interest to a wide range of stakeholders including environmental, human health, farmworker, and agricultural advocates; the chemical industry; pesticide users; and members of the public interested in the sale, distribution, or use of pesticides. Since others also may be interested, the Agency has not attempted to describe all the specific entities that may be affected by this action.

B. What should I consider as I prepare my comments for EPA?

1. *Submitting CBI.* Do not submit this information to EPA through regulations.gov or email. Clearly mark the part or all of the information that you claim to be CBI. For CBI information in a disk or CD–ROM that you mail to EPA, mark the outside of the disk or CD–ROM as CBI and then identify electronically within the disk or CD–ROM the specific information that is claimed as CBI. In addition to one complete version of the comment that includes information claimed as CBI, a copy of the comment that does not contain the information claimed as CBI must be submitted for inclusion in the public docket. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

2. *Tips for preparing your comments.* When submitting comments, remember to:

i. Identify the document by docket ID number and other identifying information (subject heading, **Federal Register** date and page number).

ii. Follow directions. The Agency may ask you to respond to specific questions or organize comments by referencing a Code of Federal Regulations (CFR) part or section number.

iii. Explain why you agree or disagree; suggest alternatives and substitute language for your requested changes.

iv. Describe any assumptions and provide any technical information and/or data that you used.

v. If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.

vi. Provide specific examples to illustrate your concerns and suggest alternatives.

vii. Explain your views as clearly as possible, avoiding the use of profanity or personal threats.

viii. Make sure to submit your comments by the comment period deadline identified.

3. *Environmental justice.* EPA seeks to achieve environmental justice, the fair treatment and meaningful involvement of any group, including minority and/or low income populations, in the development, implementation, and enforcement of environmental laws, regulations, and policies. To help address potential environmental justice issues, the Agency seeks information on any groups or segments of the population who, as a result of their location, cultural practices, or other factors, may have atypical or disproportionately high and adverse human health impacts or environmental effects from exposure to the pesticide(s) discussed in this document, compared to the general population.

II. Authority

EPA is initiating its reviews of the pesticides identified in this document pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide

Act (FIFRA) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. Section 3(g) of FIFRA provides, among other things, that the registrations of pesticides are to be reviewed every 15 years. Under FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

III. Registration Reviews

A. What action is the Agency taking?

As directed by FIFRA section 3(g), EPA is reviewing the pesticide registrations identified in the table in this unit to assure that they continue to satisfy the FIFRA standard for registration—that is, they can still be used without unreasonable adverse effects on human health or the environment. A pesticide's registration review begins when the Agency establishes a docket for the pesticide's registration review case and opens the docket for public review and comment. At present, EPA is opening registration review dockets for the cases identified in the following table.

TABLE—REGISTRATION REVIEW DOCKETS OPENING

Registration review case name and No.	Docket ID No.	Chemical review manager, telephone number, email address
Abamectin (Case 7430)	EPA–HQ–OPP–2013–0360	Khue Nguyen, (703) 347–0248, *nguyen.khue@epa.gov.*
Ametryn (Case 7036)	EPA–HQ–OPP–2013–0249	Molly Clayton, (703) 603–0522, *clayton.molly@epa.gov.*
Atrazine (Case 0062)	EPA–HQ–OPP–2013–0266	Monica Wait, (703) 347–8019, *wait.monica@epa.gov.*
Captan (Case 0120)	EPA–HQ–OPP–2013–0296	Christina Scheltema, (703) 308–2201, *scheltema.christina@epa.gov.*
Coniothyrium minitans strain CON/M/91–08 (Case 6022)	EPA–HQ–OPP–2013–0259	Jeannine Kausch, (703) 347–8920, *kausch.jeannine@epa.gov.*
Fenhexamid (Case 7027)	EPA–HQ–OPP–2013–0187	Joel Wolf, (703) 347–0228, *wolf.joel@epa.gov.*
Halohydantoins (Case 3055)	EPA–HQ–OPP–2013–0220	Sandra O' Neill, (703) 347–0141, *oneill.sandra@epa.gov.*
Indoxacarb (Case 7613)	EPA–HQ–OPP–2013–0367	Katie Weyrauch, (703) 308–0166, *weyrauch.katie@epa.gov.*
Meat Meal (Case 6041)	EPA–HQ–OPP–2013–0361	Leonard Cole, (703) 305–5412, *cole.leonard@epa.gov.*
Mesosulfuron-methyl (Case 7277)	EPA–HQ–OPP–2012–0833	Jolene Trujillo, (703) 347–0103, *trujillo.jolene@epa.gov.*
Methoxyfenozide (Case 7431)	EPA–HQ–OPP–2012–0663	Bonnie Adler, (703) 308–8523, *adler.bonnie@epa.gov.*
Metofluthrin (Case 7445)	EPA–HQ–OPP–2012–0105	Veronica Dutch, (703) 308–8585, *dutch.veronica@epa.gov.*
Propazine (Case 7278)	EPA–HQ–OPP–2013–0250	Molly Clayton, (703) 603–0522, *clayton.molly@epa.gov.*
Propylene and dipropylene glycol (Case 3126)	EPA–HQ–OPP–2013–0218	Elizabeth Hernandez, (703) 347–0241, *hernandez.elizabeth@epa.gov.*
Pymetrozine (Case 7474)	EPA–HQ–OPP–2013–0368	Steven Snyderman (703) 347–0249, *snyderman.steven@epa.gov.*
Simazine (Case 7280)	EPA–HQ–OPP–2013–0251	Molly Clayton, (703) 603–0522, *clayton.molly@epa.gov.*
Tralopyril (Case 5144)	EPA–HQ–OPP–2013–0217	Wanda Henson, (703) 308–6345, *henson.wanda@epa.gov.*
Triethylene glycol (Case 3146)	EPA–HQ–OPP–2013–0219	Elizabeth Hernandez, (703) 347–0241, *hernandez.elizabeth@epa.gov.*
Trifloxysulfuron-sodium (Case 7260)	EPA–HQ–OPP–2013–0409	Kelly Ballard, (703) 305–8126, *ballard.kelly@epa.gov.*

This document also announces the registration review case closures for the pesticides ODM (case # 0258) and resmethrin (case # 0421), and the availability of their respective Case Closure Documents. For ODM, the Notice of Receipt of a Request to Voluntarily Cancel Certain Pesticide Registrations was issued on February 20, 2013, for a 30 day comment period (78 FR 11881) (FRL–9378–9); the Agency did not receive any comments. On May 1, 2013, the Agency published the Cancellation Order for all ODM product registrations in the **Federal Register** (78 FR 25438) (FRL–9384–7). Due to the publication of the Cancellation Order for all registered ODM products in the United States, the Agency closed the registration review case for ODM, pursuant to 40 CFR 155.42(c). In addition to the registration review Case Closure Document, the registration review docket (EPA–HQ–OPP–2008–0328) for ODM also includes other relevant documents related to the registration review of this case. This action is not open for public comment.

For resmethrin, the Notice of Receipt of Requests to Voluntarily Cancel Certain Pesticide Registrations was published in the **Federal Register** on November 28, 2012 for a 180-day public comment period (77 FR 70998) (FRL–9370–3). This notice announced voluntary cancellation requests that would terminate the last resmethrin products registered for use in the United States. No comments that impacted the Agency's decision to grant the cancellation requests were received during the 180-day public comment period. On June 26, 2013, the Agency published the Cancellation Order for all resmethrin registrations. Due to the publication of the Cancellation Order for all registered resmethrin products in the United States, the Agency has closed the registration review case for resmethrin, pursuant to 40 CFR 155.42(c). The resmethrin registration review Case Closure Document, along with other documents relevant to the registration review of resmethrin, is available in the resmethrin registration review docket (EPA–HQ–OPP–2012–0414). This action is not open for public comment.

B. Docket Content

1. *Review dockets.* The registration review dockets contain information that the Agency may consider in the course of the registration review. The Agency may include information from its files

including, but not limited to, the following information:

- An overview of the registration review case status.
- A list of current product registrations and registrants.
- **Federal Register** notices regarding any pending registration actions.
- **Federal Register** notices regarding current or pending tolerances.
- Risk assessments.
- Bibliographies concerning current registrations.
- Summaries of incident data.
- Any other pertinent data or information.

Each docket contains a document summarizing what the Agency currently knows about the pesticide case and a preliminary work plan for anticipated data and assessment needs. Additional documents provide more detailed information. During this public comment period, the Agency is asking that interested persons identify any additional information they believe the Agency should consider during the registration reviews of these pesticides. The Agency identifies in each docket the areas where public comment is specifically requested, though comment in any area is welcome.

2. *Other related information.* More information on these cases, including the active ingredients for each case, may be located in the registration review schedule on the Agency's Web site at *http://www.epa.gov/oppsrrd1/registration_review/schedule.htm.* Information on the Agency's registration review program and its implementing regulation may be seen at *http://www.epa.gov/oppsrrd1/registration_review.*

3. *Information submission requirements.* Anyone may submit data or information in response to this document. To be considered during a pesticide's registration review, the submitted data or information must meet the following requirements:

- To ensure that EPA will consider data or information submitted, interested persons must submit the data or information during the comment period. The Agency may, at its discretion, consider data or information submitted at a later date.
- The data or information submitted must be presented in a legible and useable form. For example, an English translation must accompany any material that is not in English and a written transcript must accompany any information submitted as an audiographic or videographic record. Written material may be submitted in paper or electronic form.
- Submitters must clearly identify the source of any submitted data or information.
- Submitters may request the Agency to reconsider data or information that the Agency rejected in a previous review. However, submitters must explain why they believe the Agency should reconsider the data or information in the pesticide's registration review.

As provided in 40 CFR 155.58, the registration review docket for each pesticide case will remain publicly accessible through the duration of the registration review process; that is, until all actions required in the final decision on the registration review case have been completed.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: June 18, 2013.

Michael Goodis,

Acting Director, Pesticide Re-Evaluation Division, Office of Pesticide Programs.

[FR Doc. 2013–15325 Filed 6–25–13; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

Information Collection Being Reviewed by the Federal Communications Commission

AGENCY: Federal Communications Commission.

ACTION: Notice and request for comments.

SUMMARY: The Federal Communications Commission (FCC), as part of its continuing effort to reduce paperwork burdens, invites the general public and other Federal agencies to take this opportunity to comment on the following information collection, as required by the Paperwork Reduction Act (PRA) of 1995. Comments are requested concerning whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid Office of Management and Budget (OMB) control number.

DATES: Written PRA comments should be submitted on or before August 26, 2013. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Cathy Williams, FCC, via email *PRA@fcc.gov* and to *Cathy.Williams@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Cathy Williams at (202) 418–2918.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–1078.

Title: Rules and Regulations Implementing the Controlling the Assault of Non-Solicited Pornography and Marketing Act of 2003, CG Docket No. 04–53.

Form Number: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for-profit entities; Not-for-profit institutions; Individuals or households.

Number of Respondents and Responses: 5,443,062 respondents; 5,443,062 responses.

Estimated Time per Response: 1–10 hours (average per response).

Frequency of Response: Recordkeeping requirement; On occasion reporting requirements; Third party disclosure requirement.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this information collection is the CAN–SPAM Act of 2003, 15 U.S.C. 7701–7713, Public Law 108–187, 117 Stat. 2719.

Total Annual Burden: 30,254,373 hours.

Total Annual Cost: $16,244,026.

Nature and Extent of Confidentiality: Confidentiality is an issue to the extent that individuals and households provide personally identifiable information, which is covered under the FCC's system of records notice (SORN), FCC/CGB–1, ''Informal Complaints and Inquiries.'' As required by the Privacy Act, 5 U.S.C. 552a, the Commission also published SORN, FCC/CGB–1,